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Effectiveness of Absorber Intercooling for CO₂ Absorption from Natural Gas Fired Flue Gases Using Monoethanolamine Solvent

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Abstract

Chemical absorption using aqueous amine is one of the most feasible options for post-combustion CO₂ capture. One of the main challenges of this technology is its high energy requirements. Absorber intercooling was considered as a viable method to offer benefits in terms of solvent absorption capacity and mass transfer efficiency in CO₂ absorption processes. However, the effectiveness of absorber intercooling on overall energy requirements depends on other factors such as lean loading and liquid to gas ratio. This study evaluates the benefits of using two different configurations of absorber intercooling, i.e. “in-and-out” and “recycled” intercooling when using 30 wt % aqueous monoethanolamine (MEA) to capture 90% CO₂ from a natural gas fired turbine with 4 mol % CO₂. The Lean CO₂ loading was varied from 0.15 to 0.42 (mol CO₂/mol MEA) to determine the lean loading at which the application of intercooling is most significant. Absorber intercooling provides the most benefit at lean loading from 0.30 to 0.34. The use of in-and-out and recycle intercooling at 0.34 lean loading, provided 15.6 and 15.8 % reduction in the total equivalent work associated with 32.0 % and 36.6 % reduction in required packing area when using 1.2 times the minimum liquid flow rate. At lean loading greater than 0.34, the benefit of absorber intercooling is a trade-off between reduction of solvent flow rate and total energy requirement and the drawback of greater packing area in the absorber. The greatest saving in total equivalent work, 17%, was observed at the 0.36 lean loading associated with nearly 60% more packing area when using 1.2 times the minimum solvent flow rate. At very low lean loading and very high lean loading absorber intercooling does not offer significant benefit.

Keyword: Post-Combustion CO₂ Capture, Absorber Intercooling, Energy Efficiency, MEA, In-and-Out Intercooling, Recycled Intercooling,

1. Introduction

CO₂ emissions contribute substantially to global warming. According to the International Energy Agency (1) approximately one third of all CO₂ emissions is the result of fossil fuels combustion to generate electricity. Therefore, the interest in employing techniques to reduce CO₂ emissions from power plants has progressively risen over the past years. Post-combustion CO₂ capture (PCC) from

36 fossil fuel power plants by reactive absorption using amine solvents is the most promising and
37 attractive route, especially since it can be retrofitted existing power plants. The most widely used
38 solvent for chemical absorption is the aqueous solution of 30 wt % monoethanolamine (MEA) (1,2).
39 However, one major disadvantage of this process is its large energy requirement for solvent
40 regeneration. The energy requirement is usually provided by the power plant as steam and electricity,
41 which results in the considerable efficiency loss of the power plant. The addition of an amine-based
42 PCC plant to a natural gas combined cycle power plant leads to a power plant efficiency penalty of 7-
43 11% (3,4). Various alternative process configurations have been proposed to reduce the energy
44 requirements of such processes (5-13).

45 CO₂ capture by chemical absorption is based on a reversible reaction between CO₂ and a suitable
46 solvent. There are different approaches to save energy in such processes, such as reducing total
47 heating or cooling loads, improving temperature levels of provided coolants or heat sources, or a
48 combination of both (14). One useful method to reduce energy requirements is the application of
49 external coolers to absorber columns (14). Several studies have analysed the effectiveness of absorber
50 intercooling for post-combustion CO₂ capture (PCC) (7-10,12,13,15,16). The use of absorber
51 intercooling in petrochemical industries has proven its effectiveness in lowering overall energy
52 requirements. The effectiveness of using absorber intercooling in terms of energy consumption is
53 dependent on the absorbent and the process configuration (10).

54 For CO₂ capture, there have been a few studies investigating optimum conditions to use absorber
55 intercooling or identifying process conditions at which intercoolers will be most effective (7,8,10,13).
56 Plaza (7) thoroughly studied the application of simple absorber intercooling for 9 m MEA and 8 m
57 piperazine (PZ) for a range of lean loading with focus only on the absorber, and showed that absorber
58 intercooling is most effective at critical liquid-to-gas ratio, when the temperature bulge without
59 intercooling occurs in the middle of the column. Karimi et al. (10) studied the effectiveness of
60 absorber intercooling for MEA and diethanolamine (DEA) and showed that the best location for
61 intercooling is about one fourth to one fifth of the height of the absorber column from the bottom even
62 if the temperature bulge is closer to the top. Their results showed that the effect of absorber

63 intercooling is more pronounced for DEA especially at low lean loading, while intercooling at high
64 lean loading is better for MEA. Sachde and Rochelle (8) studied the mass transfer benefits of using
65 absorber intercooling for 90% CO₂ capture with 8 m PZ for flue gases with 4 to 27 % mole CO₂.
66 Their study concluded regardless of the flue gas CO₂ the absorber intercooling is most effective when
67 used at intermediate or mid-loading range lean loading, while at extreme loading (either low or high)
68 results showed negligible potential benefits from intercooling. In terms of CO₂ concentration, their
69 findings revealed that intercooling offers the greatest potential when used for 4% CO₂ (gas fired
70 turbine).

71 To properly evaluate the effectiveness of absorber intercooling, another parameters that will be
72 influenced by the use of intercooling are required to be evaluated as the benefits of using absorber
73 intercooling in majority of operating conditions is a trade-off between solvent rate and packing
74 requirement. Therefore, this study aims to analyse the effectiveness of two types of absorber
75 intercooling, “in-and-out” intercooling and “recycled” intercooling, when using 30 wt. % MEA to
76 remove 90% CO₂ from flue gases with approximately 4 % mole CO₂ for a range of lean loading from
77 0.15 to 0.42 (mol CO₂/mol MEA) in terms of solvent absorption capacity, absorber packing and
78 overall energy requirement. The CO₂ absorption/desorption process was modelled in Aspen Plus
79 V.8.4 to quantify the solvent flow rate, absorber packing volume, and solvent regeneration energy
80 with and without intercooling for a given lean loading. At each lean loading the optimum location of
81 absorber intercooling was identified by optimising the distribution of absorber packing. The
82 equivalent work concept was used to determine the amount of energy savings with absorber
83 intercooling. Finally the range of lean loading at which the application of absorber intercooling for 30
84 wt. % MEA is promising was identified, and the lean loading at which the highest savings were
85 obtained was defined.

86 **2. Modelling Framework**

87 The Aspen Plus[®] RateSep[™] model, with capabilities to rigorously model rate-based separations, was
88 used to simulate the absorber and stripper. The model used in Aspen Plus for the thermodynamic

89 properties is based on the work by Zhang et al. (17). The model uses the asymmetric electrolyte non-
 90 random-two-liquid (e-NRTL) property method to describe the CO₂-H₂O-MEA chemistry in liquid
 91 phase, and the Redlich-Kwong (RK) equation of state for the vapour phase. The model has been
 92 validated by Zhang et al. (17) against experimental data available in open literature. In the absorber,
 93 the reactions that involve CO₂ were described with a kinetic model. In this model, packed columns
 94 were divided into 40 identical segments (stages). For each stage, phase equilibrium, the energy and
 95 material balances, heat and mass transfer, and summation equations were
 96 determined. Effective interfacial area and liquid side mass transfer coefficients in the absorber
 97 column were determined using Bravo-Rocha-Fair correlation for structured packing. An aqueous
 98 solution of 30 wt % MEA was used with its proven robustness and popularity in industrial amine
 99 scrubbing because of its low cost per mole of amine, high heat of absorption, high rate of reaction and
 100 high absorption capacity. The segment model adopted for the absorber simulations was RateSep
 101 VPlug flow model, assuming the liquid phase bulk properties in each stage is similar to conditions at
 102 which the liquid phase leaves that stage, and the vapour phase bulk properties are the average of the
 103 inlet and outlet properties (18). The stripper reboiler section and the absorber intercooler heat
 104 exchanger were modelled as equilibrium stages with no reactions involved.

105 3. Evaluation Methodology

106 Absorber intercooling was evaluated at lean loading from 0.15 to 0.42. The absorber and stripper were
 107 modelled using structured packing and cylindrical columns. Unless otherwise stated, the packing was
 108 assumed to be Mellapak 250Y (19). Absorber simulation with and without intercooling was
 109 performed at flue gas conditions presented by Rezazadeh *et al.* (20) for 650 MW gas fired combined
 110 cycle power plant as presented in Table 1.

111 Table 1. Flue Gas Composition

Component	Composition (mole %)
N ₂	74.39
O ₂	12.37
CO ₂	3.905
H ₂ O	8.434
Ar	0.8952

112 The stripper packed height was over-specified at 20 m, resulting in a pinch in all cases. Noting that a
113 practical design of the stripper column would use an optimised packing height, over-specification of
114 the stripper packed height in this study confirms the packing was being equally utilised in all cases,
115 without additional height optimization criteria, while each case approached equilibrium, and therefore
116 providing an appropriate estimate for the energy requirement. To retain a constant compression work,
117 the stripper operating pressure was kept constant at 170 kPa (1.7 bar) in all load cases.

118 For an intercooled absorber column, there are three degrees of freedom for optimisation: lean loading,
119 liquid-to-gas (L/G) ratio, and the absorber packing volume. Lean loading and therefore the L/G ratio
120 were varied while maintaining the CO₂ removal constant. Furthermore, at each lean loading the
121 absorber packing volume was minimized by varying the height of the packing sections above and
122 below the intercooling. Results were normalized by the moles of CO₂ removed. Lean solvent and flue
123 gas inlet temperatures were 40°C in all cases. The absorber column diameter was calculated to
124 provide a 75% approach to flooding, and the column height was determined to satisfy 90% CO₂
125 removal in all cases. Benefits of two different types of intercooling were investigated: “in-and-out”
126 intercooling (simple intercooling) and “recycled” intercooling (advanced intercooling).

127 Process flow diagrams (PFD) of an absorber column with simple and advanced intercooling are
128 shown in Figures 1 and 2, respectively. In simple intercooling the semi-rich solvent exits the absorber
129 column at the end of one packing section and passes through an external heat exchanger (cooler) to
130 cool down to the temperature at which the lean solvent first enters the absorber column at the top, and
131 then returns to the column at the top of the successive packing section.

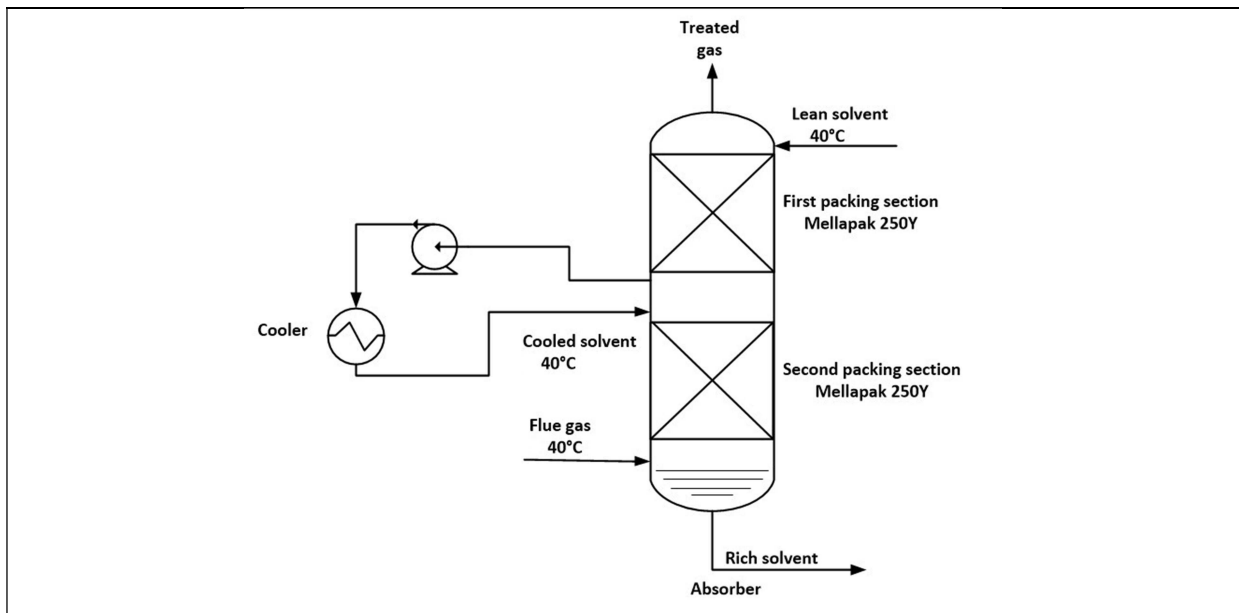


Figure 1. The arrangement of an absorber column with in-and-out (simple) intercooler

132 With advanced intercooling, the semi-rich solvent is extracted below a middle section of packing,
 133 cools in an external cooler to the temperature at which the lean solvent first entered the absorber
 134 column at the top, and returns back to the column at the top of the middle section. In this
 135 configuration, the absorber packed column was divided into three sections, by which the first and
 136 third sections were packed with the Sulzer Mellapak 250Y structured packing, and the middle section
 137 (recycled section) with a coarse structured packing, Sulzer Mellapak 125Y, to avoid excessive
 138 pressure drop due to the high solvent load in in the middle section. In essence, this is a modification of
 139 simple intercooling where the cooled semi-rich solvent recycles around the middle section. The
 140 recycle rate is usually 2 to 5 times the solvent flow rate (8) which can be optimised with respect to the
 141 operational costs for running the recycled pumps and the absorber flooding.

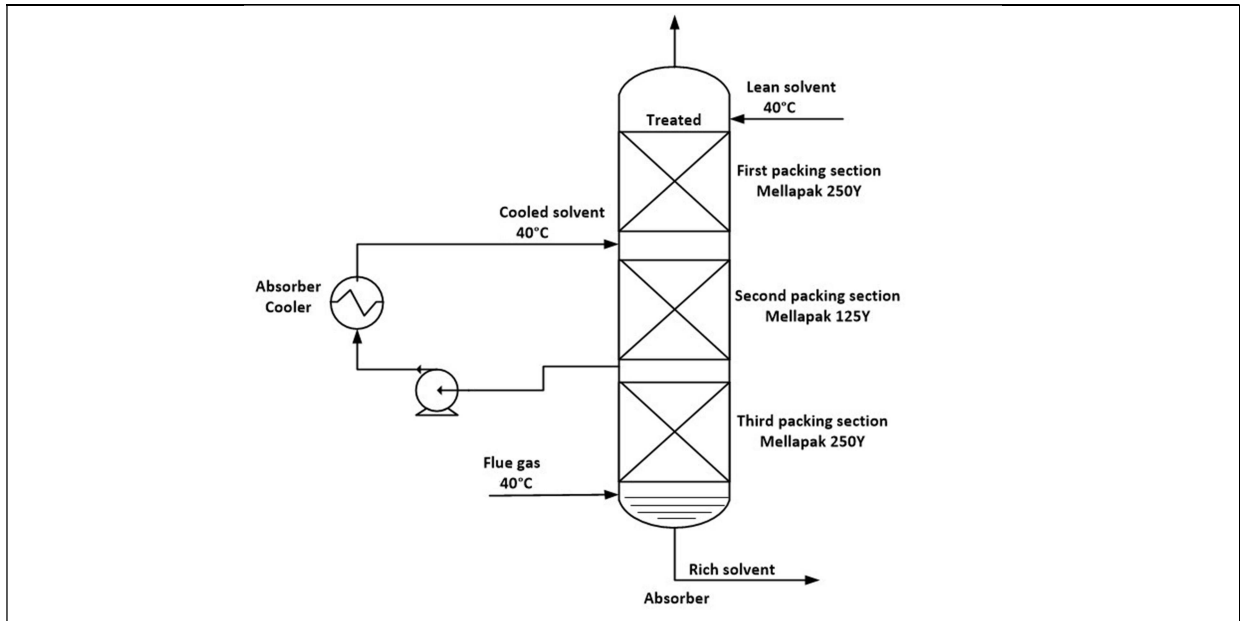


Figure 2. The arrangement of an absorber column with recycled (advanced) intercooling

142 To find a proper recycle ratio, various recycle ratios, from 1 to 9 times the solvent flow rate, were
 143 compared with each other and with the base case, a simple absorber with no intercooling (no recycle
 144 rate). As shown in Figure 3 the recycle ratio of 3 was selected as the optimum ratio for natural gas
 145 applications with 30 wt % MEA .

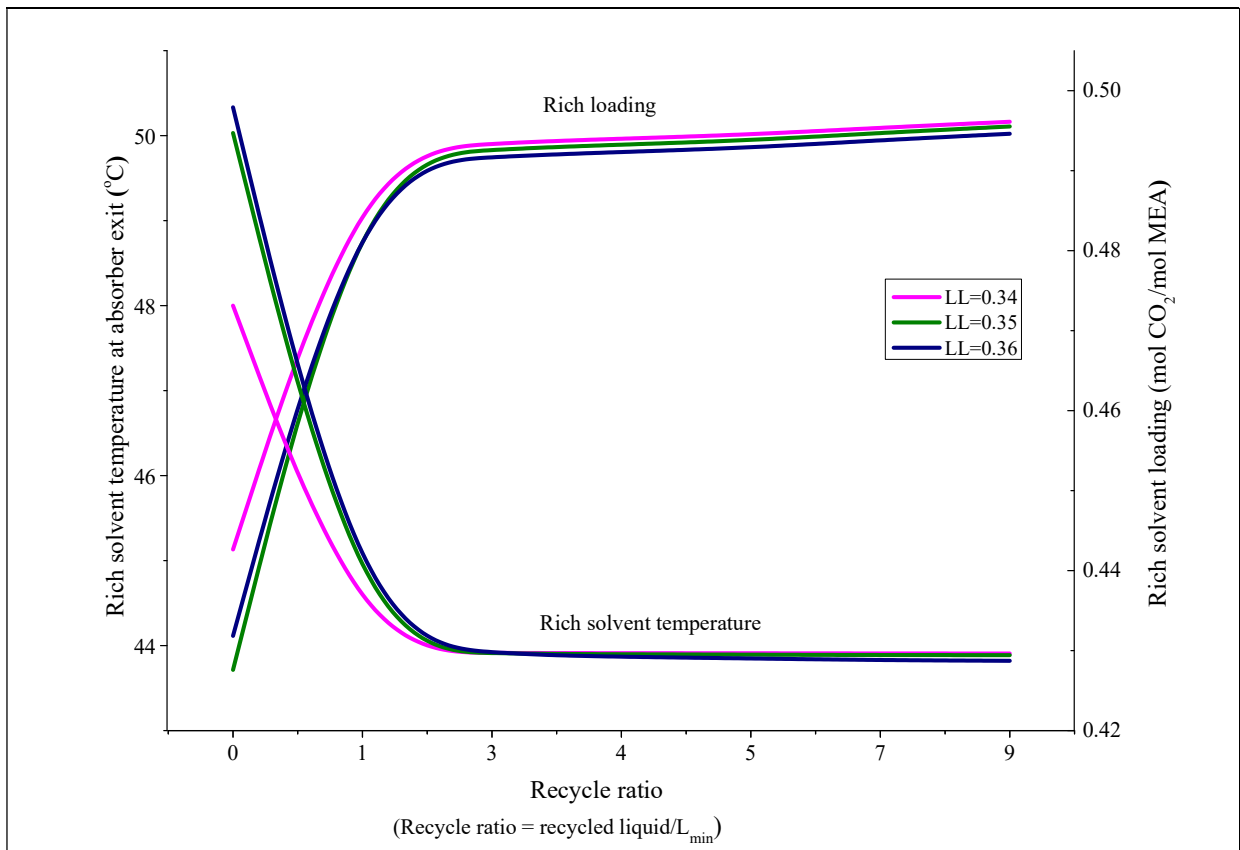


Figure 3. Variation of rich solvent loading and temperature at absorber exit with cooling solvent recycle rate

146 3.1. Overall energy requirement

147 Total equivalent work was used to evaluate the overall energy requirement with and without absorber
148 intercooling. This value estimates the total electrical work penalty from the power plant by operating
149 the stripper, compressors and pumps. The total equivalent work (W_{eq}) is calculated as the sum of the
150 regeneration heat equivalent work (W_{heat}), compression work (W_{Comp}), and pump work (W_{Pump}), as
151 shown in Eq. (1) (21).

$$W_{eq} = W_{heat} + W_{comp} + W_{pump} \quad (1)$$

152 The regeneration heat would draw steam from the steam turbine of the power plant that would be
153 otherwise expanded in low pressure steam turbines to generate electricity (22). Oyenekan (21)
154 suggested calculating the equivalent electrical penalty (work) associated with the heat required for
155 solvent regeneration using the Carnot efficiency, as expressed in Eq. (2).

$$W_{heat} = \eta_{effective} \left(\frac{T_{reb} + \Delta T - T_{sink}}{T_{reb} + \Delta T} \right) Q_{reb} \quad (2)$$

156 Where, $\eta_{effective}$ is the turbine effective efficiency, T_{reb} is the solvent temperature at the reboiler, ΔT
157 is the temperature difference between hot and cold streams at the reboiler, T_{sink} is the cooling water
158 temperature, and Q_{reb} is the reboiler heat duty. Assumptions made for Eq. 2 include a 90 % efficiency
159 to account for non-ideal expansion in steam turbines (23), an approach temperature of 5 °C for the
160 steam side in the reboiler section, and a sink temperature of 40 °C.

161 The compression work is the work to compress the captured CO₂ from the stripper pressure (P_{in}), to
162 the storage pressure, e.g. 15 MPa (150 bar), and was calculated using Eq. (3) (24,25).

$$W_{comp} = -3.48 \ln(P_{in}) + 14.85, \quad 1 < P_{in} (bar) < 20 \quad (3)$$

163 Assumptions made for Eq. (3) include a compression ratio of 2 or less for each compression stage, a
164 compressor polytropic efficiency of 86 %, inter-stage cooling to 40 °C with knocked out water
165 between stages with zero pressure drop (24).

166 For the absorber with no intercooling, the pump work includes the required head at the efficiency of
167 the pump, 75%, to move and circulate the solvent from absorber to the pressure of stripper and vice
168 versa. For the absorber with simple and advanced intercooling, the work required to pump the cooling
169 solvent from the absorber to the external cooler and back to the column is added to the pump work.
170 The flue gas blower work is excluded. The Aspen Plus pump block is used to calculate the pump
171 work.

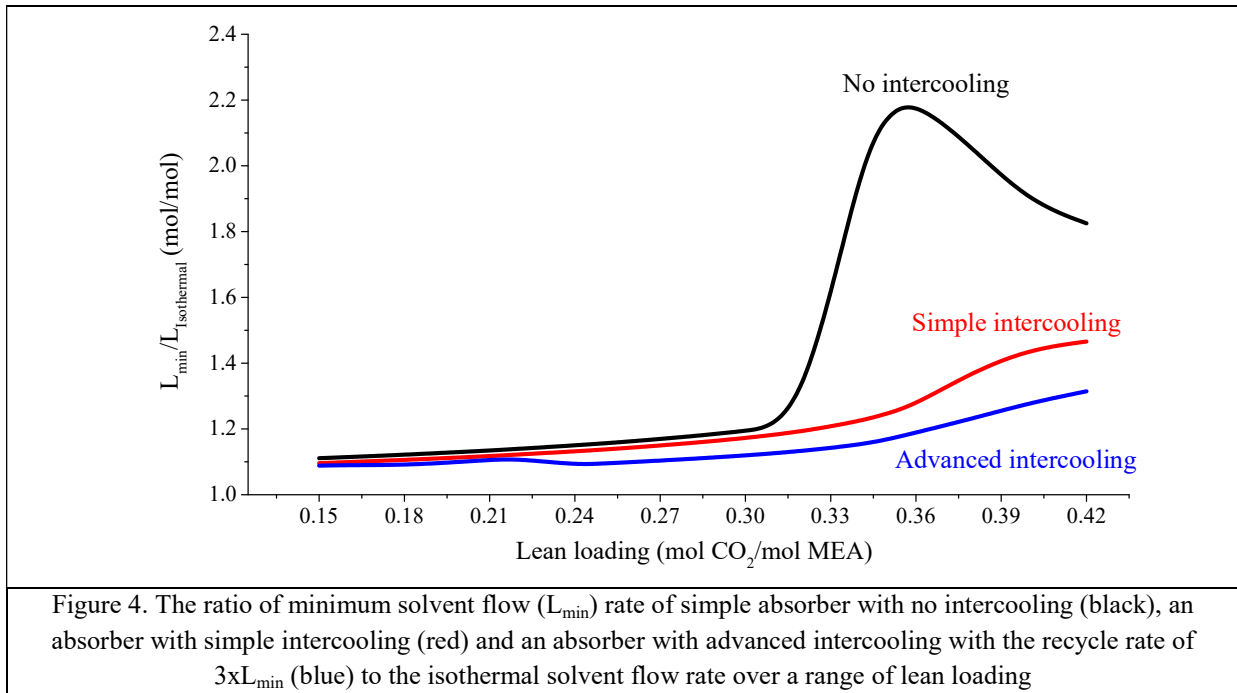
172 4. Results and discussion

173 4.1. The effect of absorber intercooling on minimum solvent flow rate (L_{\min})

174 For a given lean loading and CO₂ removal, the solvent flow is a function of packing area. By
175 increasing the packing area, the liquid flow decreases until it reaches its minimum value. Figure 4
176 shows L_{\min} to achieve 90% CO₂ removal for a range of lean loading from 0.15 to 0.42 for the three
177 absorber cases: (1) no intercooling, (2) simple intercooling, and (3) advanced intercooling. With no
178 intercooling, L_{\min} was determined with 40 m of absorber packing to assure equilibrium pinch at the
179 rich end of the column (26), provided the fractional approach to flooding was held at 75%. Similarly,
180 for absorbers with simple and advanced intercooling, for a given lean loading, L_{\min} to achieve 90%
181 CO₂ removal was determined 30 m of packing in each section with 75% flooding fraction.

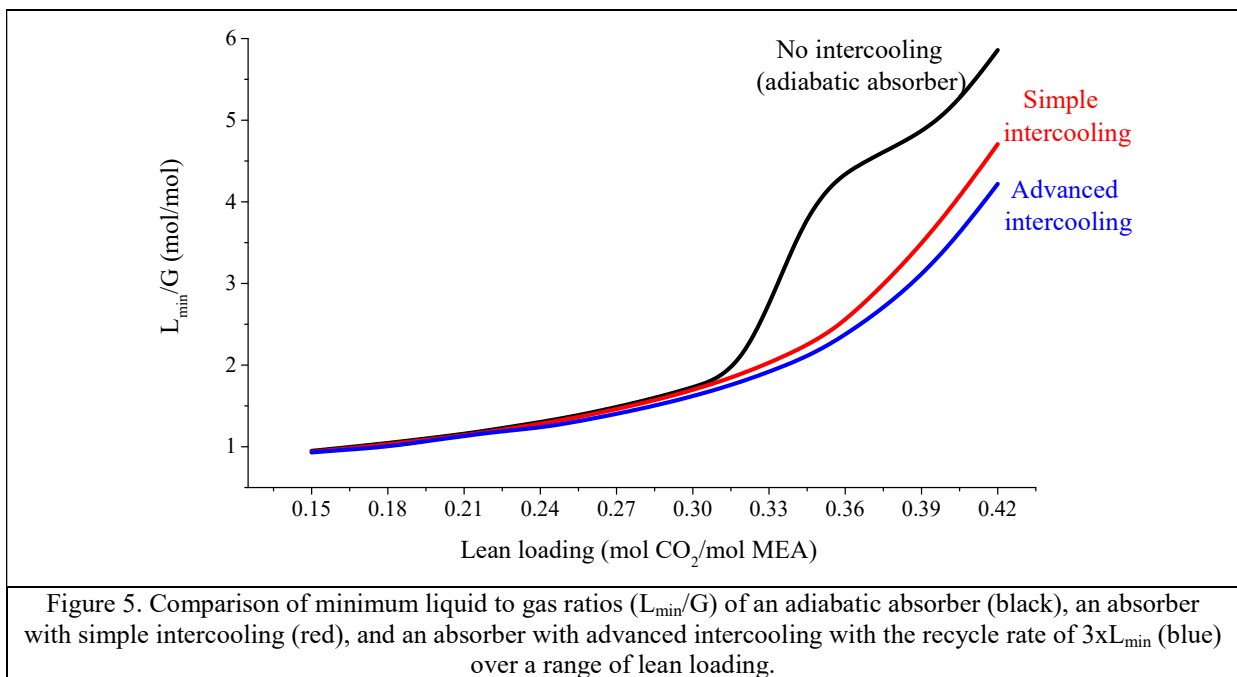
182 The effectiveness of intercooling can be better realised by comparing L_{\min} at any given lean loading in
183 relation to the theoretical minimum solvent flow rate required at that lean loading to attain 90% CO₂
184 removal rate. The theoretical minimum solvent flow rate ($L_{\text{isothermal}}$) was determined assuming an
185 isothermal absorber where the temperature of the liquid phase throughout the column is the same and
186 equal to the inlet liquid temperature (ideal intercooling) (26). As shown in Figure 4, the lean loading
187 range at which the application of intercooling is promising is equal to and higher than 0.30 as the
188 $L_{\min}/L_{\text{isothermal}}$ ratio increases by increasing lean loading. The $L_{\min}/L_{\text{isothermal}}$ ratio at lean loading below
189 0.30 is close to one, so absorber intercooling would not be helpful in this range. This figure also
190 indicates the minimum ratio is related to the advanced intercooling option suggesting its better
191 performance compared to the simple intercooling option. The highest reduction in the minimum

192 solvent flow rate offered by the simple and advanced intercooling were observed at lean loading of
 193 0.35 with 42.4% and 46.1% reduction when compared to the non-intercooled option, respectively.



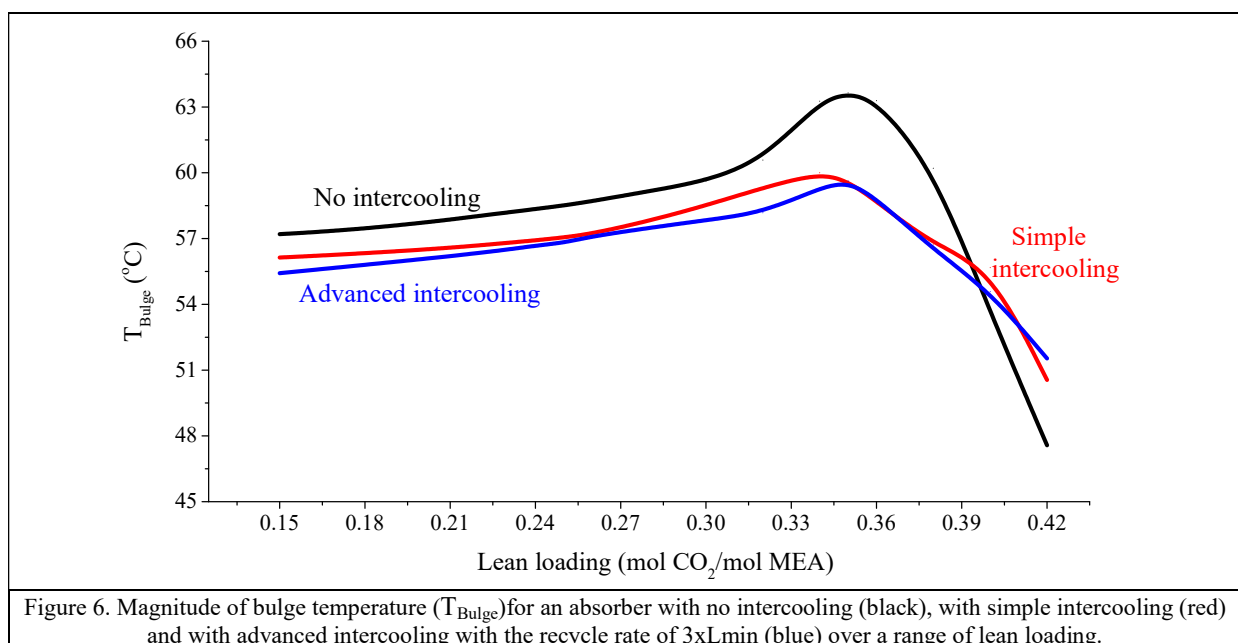
194

195 Figure 5 compares L_{\min}/G with no intercooling, simple intercooling, and advanced intercooling.



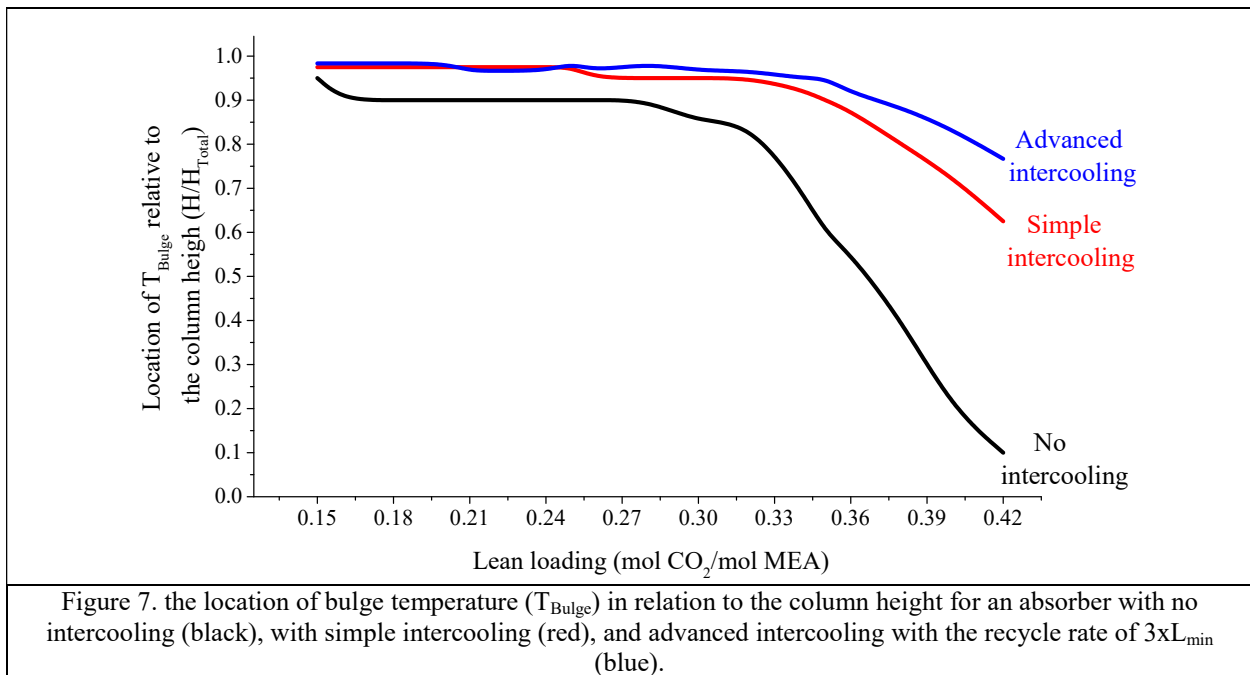
196 **4.2. The effect of absorber intercooling on temperature bulge**

197 The aqueous solvent enters the absorber column at the top and counter-currently contacts the flue gas.
 198 As the solvent absorbs the CO₂, its temperature increases and causes water to vaporise. Toward the
 199 top of the column, the produced water vapour condenses by contacting counter-currently the cooler
 200 solvent, which leads to formation of a pronounced temperature bulge in the gas and liquid temperature
 201 profiles (27). The magnitude and location of the temperature bulge depends on the solvent lean
 202 loading and L/G. Figure 6 shows the magnitude of bulge temperature (T_{Bulge}) for a range of lean
 203 loading for an absorber with no intercooling, with simple intercooling, and with advanced
 204 intercooling.



205 Figure 7 shows the location of bulge in relation to the absorber column height. As L/G increases, the
 206 location of the bulge moves toward the bottom of the column and its magnitude decreases as more
 207 heat has been carried by the solvent due to its relatively higher heat capacity. As shown in Figure 6, at
 208 low lean loading ($0.15 < \text{lean loading} < 0.30$), the bulge occurs at the top of the packed column. As
 209 lean loading and therefore L/G increases, the location of the bulge moves toward the bottom of the
 210 column. The slope of move is more pronounced for the absorber with no intercooler. Concurrently
 211 the magnitude of bulge temperature ascends by which the greatest temperature bulge occurred at the
 212 lean loading of 0.35 in all three cases. After this point, as lean loading increases, the magnitude of
 213 temperature bulge descends. The temperature bulge at its peak is located near the middle of the
 214 column ($H_{\text{Bulge}}/H_{\text{total}}=0.6$) in an absorber with no intercooling, while for the absorber with simple

215 intercooling and advanced intercooling, the temperature bulge at its peak occurs near the top of the
 216 packed column, ($H_{\text{Bulge}}/H_{\text{total}}=0.925$) and ($H_{\text{Bulge}}/H_{\text{total}}=0.95$), respectively.

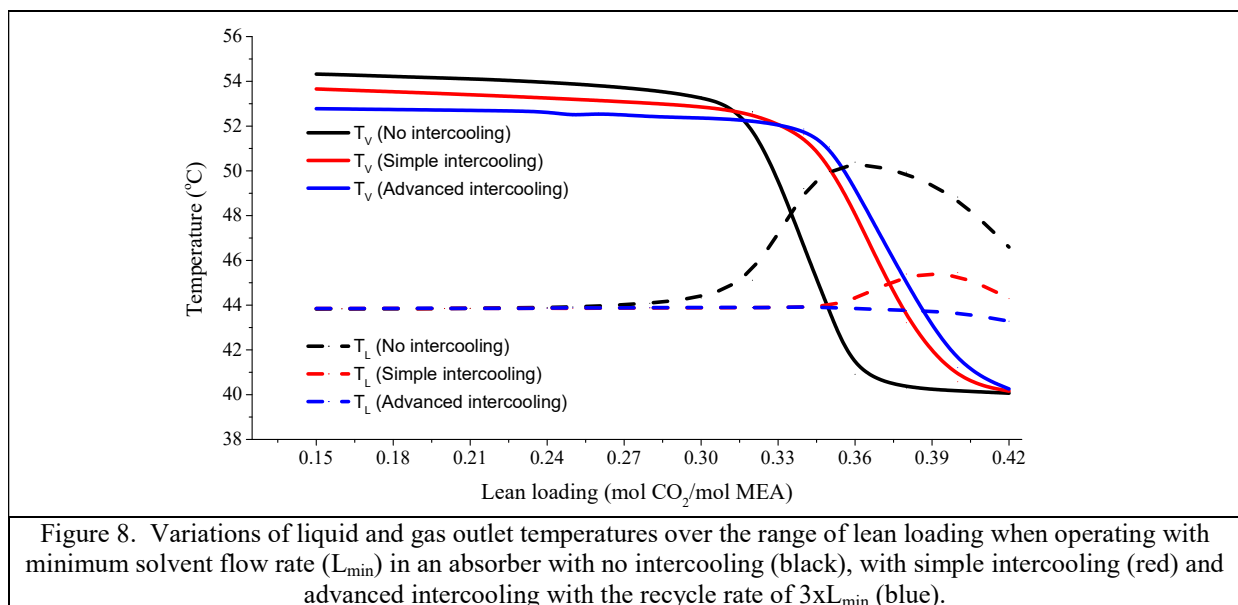


217 Absorber performance is set by keeping the rich solvent fully saturated and using the solvent flow rate
 218 to maintain the desired removal rate. Such an absorber is called rich-end pinched (26). However, at
 219 higher L/G ratios, there is excess solvent relative to the inlet gas, therefore fully saturated rich solvent
 220 could not be kept, such an absorber is called lean end pinched (27). According to the T_{bulge} theory
 221 (27), the greatest absorption rate will occur away from pinch and so does the temperature bulge.
 222 Therefore, as long as the temperature bulge occurs away from the equilibrium pinch, its effect on the
 223 column mass transfer is negligible. As can be observed from Figures 5 and 7, for the absorber with no
 224 intercooling, at lean loadings between 0.32 and 0.36, the sharp rise in L/G coincides with the location
 225 of temperature bulge being near the middle of the column.

226 Curves related to simple and advanced intercooling shown in Figures 6 and 7 confirm the use of
 227 absorber intercooling changes the location and the magnitude of the temperature bulge. The maximum
 228 bulge temperature after incorporating simple and advanced intercooling dropped to 60.0°C and
 229 59.6°C respectively, compared to 63.6°C without intercooling. Concurrently, employing absorber
 230 intercooling favours the column mass transfer efficiency by moving the temperature bulge to the top
 231 of the column. The location of temperature bulge moves to 0.925 and 0.950 of the total absorber

232 packed height, when simple and advanced intercooling were applied, respectively, compared to 0.60
233 in the non-intercooled case.

234 In an absorber with no intercooling, when the temperature bulge occurs near or at the middle of the
235 packed column, it is defined as the critical temperature bulge (27) with the critical L/G. In this study,
236 the critical temperature bulge was realised at lean loading of 0.36, with critical L/G of 4.45 (mol/mol).
237 The magnitude and location of the bulge temperature at the critical lean loading are 63.3°C and
238 $H_{\text{Bulge}}/H_{\text{Total}}=0.55$, respectively. Figure 8 shows the variation of liquid (rich solvent) and gas (treated
239 solvent) temperatures when leaving the absorber column of the three cases. As shown, both liquid and
240 gas temperature curves display a smoother trend after employing absorber intercooling. The effect of
241 intercooling on the liquid outlet temperature is more pronounced especially in the advanced
242 intercooling case. This is due to the solvent having in general a cooler temperature profile along the
243 absorber column after employing intercooling, which results in an increase in the solvent absorption
244 capacity since the absorption capacity of amine solvents for CO₂ increases with lower temperature.
245 Equally, for a fixed CO₂ removal, absorber intercooling requires less solvent, as shown in Figure 5.

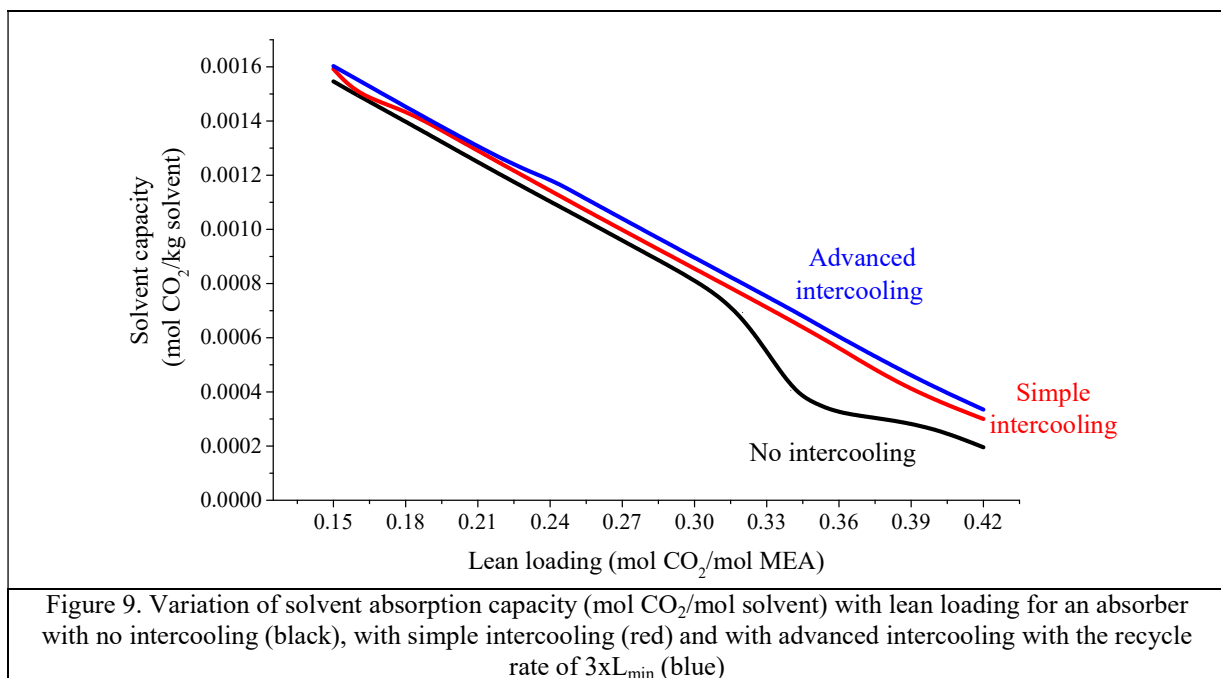


246 4.3. Effect of absorber intercooling on Solvent capacity

247 Solvent capacity to absorb CO₂ increases as temperature decreases (15). The solvent capacity is
248 defined as moles CO₂ removed per kg lean solvent. Figure 9 shows the variation of solvent absorption
249 capacity with lean loading for the three cases with 90% CO₂ removal. With no intercooling the

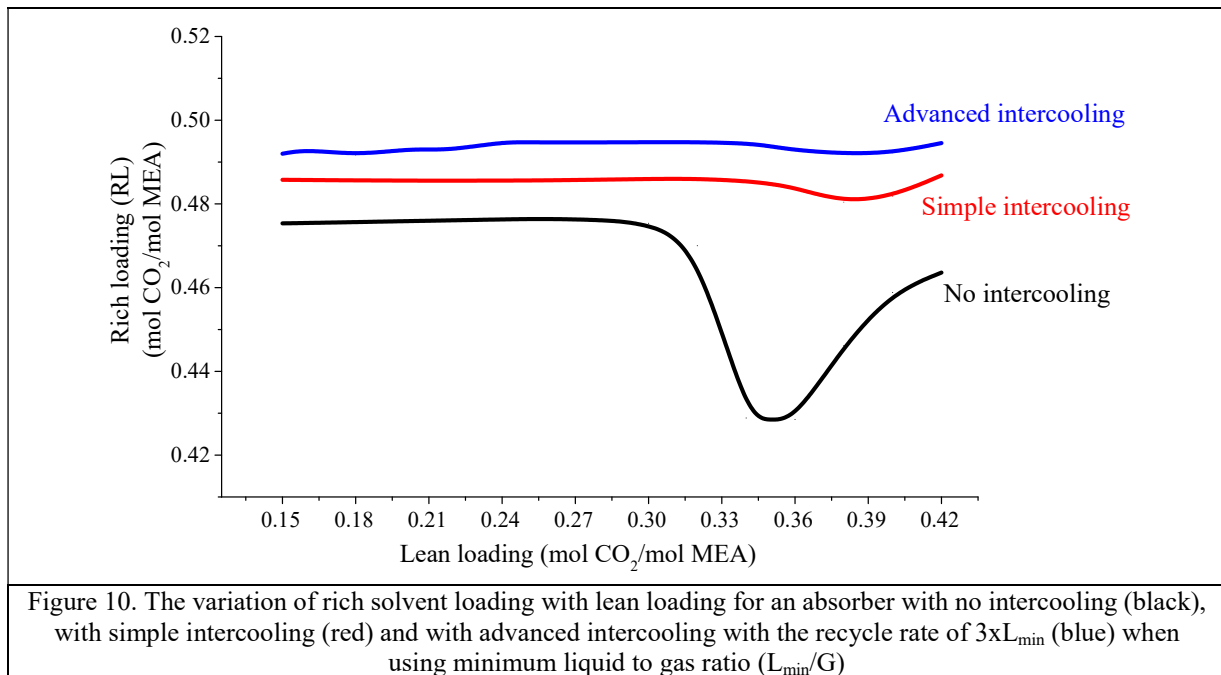
250 solvent capacity substantially decreased after a lean loading of 0.32. The rate of solvent absorption
251 capacity reduction is more pronounced from lean loading 0.32 to 0.36. Lean loading 0.36 is the
252 critical lean loading. After the critical lean loading, a slight improvement in capacity was observed
253 due to the excessive increase in the liquid to gas ratio at those lean loading as shown in Figure 5.

254 Figure 9 shows the change in the solvent capacity when using absorber intercooling. At 0.32 lean
255 loading and above the use of absorber intercooling significantly improves the solvent capacity. At
256 0.34 lean loading, the use of simple and advanced intercooling provide 75 % and 88 % increase in the
257 solvent capacity, respectively. In general, the solvent capacity decreases with increasing lean loading
258 due to the limiting capacity imposed by the initial high CO₂ content in the lean feed.



259 4.4. Effect of absorber intercooling on rich solvent loading

260 Figure 10 compares the variation of rich solvent loading with lean loading with no intercooling,
261 simple intercooling and advanced intercooling.



262 By considering Figures 5, 6 and 10 together, the following results can be concluded:

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- At lean loading up to 0.30, the rich loading with no intercooling is fairly constant with a steady increase of L_{\min}/G with increasing lean loading. Using both simple and advanced intercooling slightly increase the rich loading by 2.0% and 3.8%, respectively, with no noticeable changes in their L_{\min}/G . At this range, in all three cases the temperature bulge occurs at the top of the column confirming the use of absorber intercooling would not be helpful.
 - At lean loading from 0.30 to 0.36, a noticeable decline in the rich loading coincided with a sharp increase in the L_{\min}/G were observed at the absorber with no intercooling. At this range, the temperature bulge occurs near the middle of the column. The difference between the L_{\min}/G of non-intercooled and intercooled cases reaches its maximum at 0.34 lean loading. The significant reduction in L_{\min}/G and improvement in rich loading by using simple and advanced intercooling confirm the effectiveness of intercooling at this lean loading range. At 0.34 lean loading, the use of simple and advanced intercooling provides 42.0% and 45.6% reduction in L_{\min}/G , and 12.4 and 14.5% increase in rich loading, respectively. Also, at 0.34 lean loading, the use of simple and advanced intercooling resulted in 63.5% and 73.6% increase in the solvent absorption capacity, respectively.

279 • At lean loading higher than 0.36, a gradual increase in the rich loading coincided with
280 continual increase in the L_{\min}/G with lean loading observed at the absorber with no
281 intercooling. Due to the limited capacity, the solvent flow considerably increases with lean
282 loading. The use of absorber intercooling slightly reduces the solvent flow yet the rich loading
283 remain almost constant.

284 As shown in Figure 10, the increase in rich loading by using simple and advanced intercooling
285 confirms that intercooling in general allows the absorber column to have a closer approach to
286 equilibrium. Furthermore, for a given lean loading, the increase in rich loading coincides with another
287 advantage of using intercooling that is less lean solvent flow is required compared to that of no
288 intercooling to achieve 90% CO₂ removal. As shown, the use of absorber intercooling is helpful at
289 medium to high lean loading which is associated with higher solvent flow. The benefit of absorber
290 intercooling at high lean loading should be realised by evaluating the energy requirement for solvent
291 regeneration. This will be discussed in the following sections.

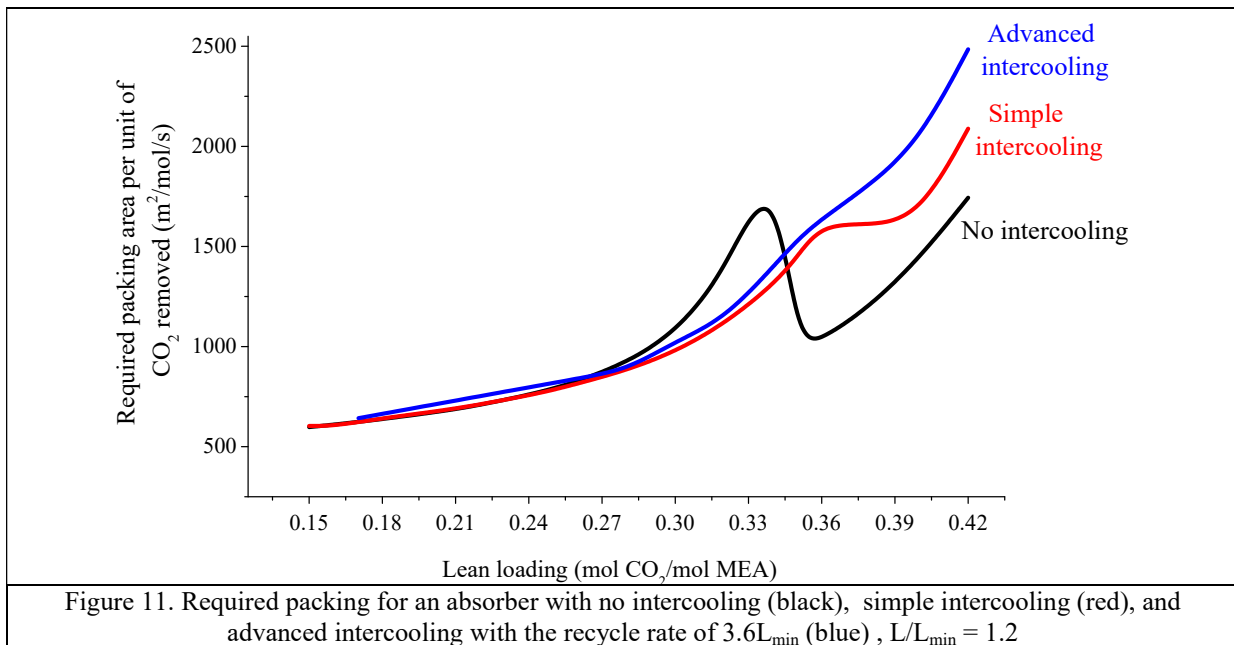
292 **4.5. Application of Absorber intercooling with 1.2 L_{\min}**

293 The lean loading range at which the use of absorber intercooling is beneficial when using minimum
294 liquid flow (L_{\min}) is roughly from 0.30 to 0.38. The minimum liquid flow to achieve 90% CO₂
295 removal is determined based on an infinite packing volume, which is not a practical design in terms of
296 plant economics. The optimisation of liquid to gas ratio in terms of plant economics suggests the
297 molar L/G ratio should be about 1.2 to 1.5 times its minimum value in order to avoid using excessive
298 packing (26). Therefore, the solvent flow was set to 1.2 times its minimum flow. Subsequently, the
299 absorber required packing volume, including each section height and diameter, was optimised to
300 provide 90% CO₂ removal.

301 **4.5.1. Effect of absorber intercooling on absorber packing area with 1.2 L_{\min}**

302 Figure 11 shows the required packing area to achieve 90% CO₂ removal when using 1.2 L_{\min} . The
303 required packing area is calculated by multiplying the volume of packing by the packing specific
304 surface area. For Sulzer Mellapak 250Y and 125Y, the specific surface area is 250 and 125 m²/m³

305 packing, respectively (19). For all cases, the optimum packing volume was calculated by with
 306 diameter specified to get 75 % flooding, and adjusting the height to achieve 90% CO₂ removal.



307 As Figure 11 shows, at lean loading from 0.28 to 0.34, the required packing area decreases when
 308 using absorber intercooling with $1.2 L_{\min}$. The greatest reduction in the required packing area was
 309 observed at 0.34 lean loading with 30% and 26% reduction when incorporating simple and advanced
 310 intercooling, respectively.

311 At 0.35 lean loading and above, the use of absorber intercooling resulted in greater rich loading. As
 312 Figure 11 shows, additional packing area is required to achieve these benefits. For instance, at 0.36
 313 lean loading, the use of simple and advanced intercooling, results in 43 % and 47 % reduction in the
 314 L/G, respectively, which is associated with 60 % and 62 % increase in the absorber required packing
 315 area, respectively. At lean loading below 0.30 the use of absorber intercooling does not change the
 316 packing requirement.

317 The absorber required packing areas per unit of CO₂ removed as presented in Figure 11 for a range of
 318 lean loading with and without using absorber intercooling were calculated based on using the Bravo-
 319 Rocha-Fair correlation to determine the liquid side mass transfer coefficient (k_L) which is the Aspen
 320 Plus® suggested mass transfer model to calculate the liquid side mass transfer coefficient for
 321 structured packing applications. There is a great deal of uncertainty in calculating the liquid side mass

322 transfer coefficient, and this uncertainty directly impacts the calculation of absorber packing area. The
323 Bravo-Rocha-Fair correlation is a generalised mass transfer model which represents an average of a
324 wide range of hydraulic conditions, packing types/materials, and fluid properties that may not be
325 representative of chemical based CO₂ capture process conditions using amine solvents (28). A new
326 empirical mass transfer model developed by Sachde (28), called the sachde model, to isolate
327 independent variables that impact mass transfer performance and to regress model coefficients from
328 data collected in a pilot scale column operated with structured packing for chemical based CO₂
329 capture process. The Sachde model is expected to closely represent the packing and hydraulic
330 conditions experienced in the amine-based absorption columns used in CO₂ capture processes. The
331 Sachde model was developed using data collected at the Separation Research Program (SRP) at the
332 University of Texas at Austin (UT) (28).

333 To put this uncertainty into perspective, for 0.36 lean loading, the liquid side mass transfer coefficient
334 when using simple and advanced intercooling, was calculated using these two models and presented
335 in Table 2.

336 Table 2. Comparison of the liquid side mass transfer coefficients (k_L) calculated using the Sachde (28) and those
337 using Bravo et al. (1985) at 0.36 lean loading

Method of determining k_L (m/s)	Simple intercooling	Advanced intercooling
Bravo-Rocha-Fair	3.51	4.76
Sachde	1.57	4.78

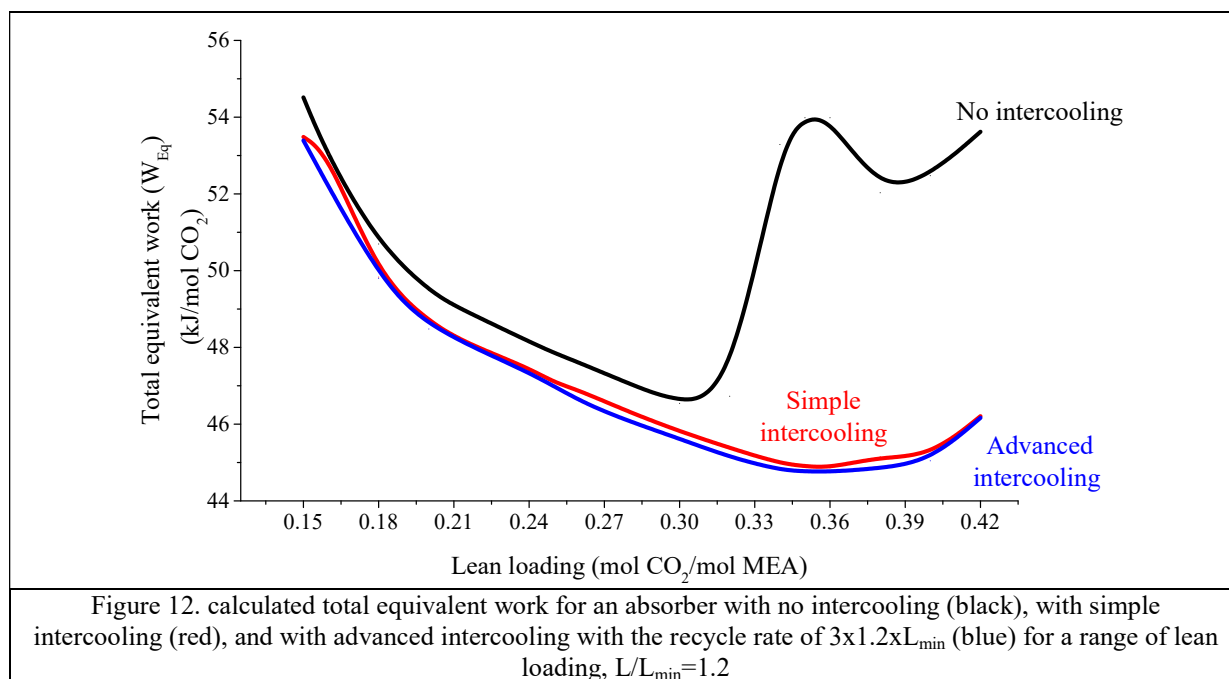
338 As shown in Table 2, the liquid side mass transfer coefficient calculated by Bravo-Rocha-Fair model
339 is more than twice that calculated by Sachde when using simple intercooler. The calculation and
340 comparison of the liquid-side mass transfer coefficient using these two approaches, as presented in
341 Table 2, were performed for a part of lean loading range that requires additional packing area to
342 realise benefits of using absorber intercooling. As such, the present work only covers a limited range
343 of process conditions, i.e. those that are potentially industrially relevant. In contrast, investigating the
344 origins of differences between these two approaches would require a very different approach, in
345 which a wide range of process conditions would be investigated, to determine the circumstances in

346 which the two approaches converge, versus those under which divergence between the two
347 approaches takes place and this would be a necessary precursor to drawing a final conclusion on
348 the origins of differences between the estimation of the liquid-side mass transfer coefficient each of
349 these two approaches provides.

350 Calculations with the Sachde correlation show that at 0.36 lean loading, the use of advanced
351 intercooling results in nearly 14.5% reduction in the required packing area compared to that when
352 simple intercooling was used, while calculations based on the Bravo-Rocha-Fair model show 1.5%
353 more packing area is required when using advanced intercooling than that of simple intercooling.

354 4.5.2. Effect of absorber intercooling on total equivalent work

355 Figure 12 shows the total equivalent work of the CO₂ capture process with and without absorber
356 intercooling over the range of lean loading. The compression work was constant across all cases as the
357 stripper pressure was kept at 17 kPa (1.7 bar).



358 As Figure 12 shows, at 0.30 lean loading and above, the total equivalent work significantly decreases
359 with absorber intercooling. The highest energy saving (17%) was realised at 0.36 lean loading with
360 both simple and advanced intercooling. Figures 5, 11, and 12 demonstrate that absorber intercooling at
361 lean loading from 0.30 to 0.34 reduces solvent flow, absorber packing area, and total equivalent work.
362 The use of simple and advanced intercooling at 0.34 lean loading decreases the total equivalent work

363 by 16% with 32 % and 37 % reduction in packing area. At lean loading from 0.30 to 0.34 absorber
364 intercooling is promising and helpful.

365 With greater than 0.35 lean loading and above, the benefits of absorber intercooling are a trade-off
366 between reduction of solvent flow and total energy requirement and the use of greater packing area in
367 the absorber.

368 **5. Conclusions**

369 Two absorber intercooling configurations were evaluated for CO₂ capture with 30 wt % MEA to
370 remove 90 % CO₂ from gas turbine fired flue gas for lean loading from 0.15 to 0.42. The effect of
371 absorber intercooling on temperature bulge, liquid flow, L/G, rich loading, and solvent capacity were
372 evaluated using minimum solvent flow (L_{min}). Benefits of using absorber intercooling on the absorber
373 packing area and the plant overall energy requirement were quantified using 1.2 L_{min} . The total
374 equivalent work value was used to evaluate the plant overall energy requirement.

375 At lean loading below 0.30, the temperature bulge occurs near the top of the column and away from
376 the equilibrium pinch at the rich-end with no intercooling, therefore absorber intercooling would not
377 be helpful in this range. Minor reduction in L_{min}/G and total equivalent work with intercooling
378 confirms this conclusion.

379 At lean loading from 0.30 and 0.36, significant increase in L_{min}/G coincides with sharp reduction in
380 rich loading with no intercooling. In this range, the temperature bulge was around the middle of the
381 column. The use of absorber intercooling showed a positive effect on both L_{min}/G and rich loading.
382 The use of absorber intercooling at lean loading from 0.30 and 0.34 provides reduction in both
383 required packing area and total equivalent work. At 0.34 lean loading, incorporation of simple and
384 advanced intercooling provides respectively 32% and 37% reduction in the required packing area,
385 coinciding with 16%% reduction in the total equivalent work.

386 At lean loading of 0.35 and above, absorber intercooling reduces L/G, rich loading, and the overall
387 energy requirement. In this range, additional packing is needed at $L/L_{min}=1.2$ to get these benefits.
388 For instance, at 0.36 lean loading, simple and advanced intercooling provide 43% and 47% reduction

389 in L/G, 17% reduction in the total equivalent work, and 60% and 62% increase in the absorber
390 packing area, respectively.

391 There is a considerable difference between the calculated value of the absorber liquid side mass
392 transfer coefficient (k_L) when using the Bravo-Rocha-Fair correlation and that determined by the
393 Sachde (28) correlation, resulting in a great difference in the estimated packing area requirement at
394 higher lean loading (0.36 and above) when using simple and advanced intercooling. At 0.36 lean
395 loading, the Sachde correlation with advanced intercooling results in 15% reduction in the required
396 packing area compared simple intercooling, while calculations with Bravo-Rocha-Fair require 1.5%
397 more packing area than with simple intercooling.

398 These findings can be used as a guideline for future applications of absorber intercooling for
399 commercial scale natural gas fired turbines with 4 mol % CO₂ when using 30 wt % aqueous MEA as
400 solvent.

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